

Ajovy® Prior Authorization Request Form (Page 1 of 2)
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Member Information (required)			Provider Information (required)		
Member Name:			Provider Name:		
Insurance ID#:			NPI#:		Specialty:
Date of Birth:			Office Phone:		
Street Address:			Office Fax:		
City:	State:	Zip:	Office Street Address:		
Phone:			City:	State:	Zip:
Medication Information (required)					
Medication Name:			Strength:		Dosage Form:
<input type="checkbox"/> Check if requesting brand			Directions for Use:		
<input type="checkbox"/> Check if request is for continuation of therapy					
Clinical Information (required)					
Select the diagnosis below: <input type="checkbox"/> Chronic migraines <input type="checkbox"/> Episodic migraines <input type="checkbox"/> Other diagnosis: _____ ICD-10 Code(s): _____					
Clinical information: Is the requested medication prescribed by or in consultation with a neurologist or pain specialist? <input type="checkbox"/> Yes <input type="checkbox"/> No Select if the patient has failed a 2-month trial of the following migraine prophylactic classes supported from The American Headache Society/American Academy of Neurology treatment guidelines: <input type="checkbox"/> Anticonvulsants (divalproex, valproate, topiramate) <input type="checkbox"/> Antidepressants (amitriptyline, venlafaxine) <input type="checkbox"/> Beta blockers (atenolol, metoprolol, nadolol, propranolol, timolol) Select if the patient has a trial and failure, contraindication, or intolerance to the following preferred agents: <input type="checkbox"/> Aimovig <input type="checkbox"/> Emgality					
Chronic migraines, also answer the following: Does the patient have ≥ 15 headache days per month AND > 8 migraine days per month for a minimum of 3 months? <input type="checkbox"/> Yes <input type="checkbox"/> No Has the patient been evaluated for medication overuse headache (MOH) (defined as headaches occurring greater than or equal to 15 days per month, and develops as a consequence of regular overuse of acute or symptomatic headache medication for more than 3 months)? <input type="checkbox"/> Yes <input type="checkbox"/> No Will the treatment include a plan to taper off the offending medication if MOH is diagnosed? <input type="checkbox"/> Yes <input type="checkbox"/> No Will the patient be initiating botulinum toxin headache prophylaxis after starting the requested agent? <input type="checkbox"/> Yes <input type="checkbox"/> No					
Episodic migraines, also answer the following: Does the patient have < 15 headache days per month AND 4 to 14 migraine days per month for a minimum of 3 months? <input type="checkbox"/> Yes <input type="checkbox"/> No					
Reauthorization: If this is a reauthorization request, answer the following: Is the requested medication prescribed by or in consultation with a neurologist or pain specialist? <input type="checkbox"/> Yes <input type="checkbox"/> No Has the patient experienced a positive response to therapy, demonstrated by a reduction in headache frequency and/or intensity? <input type="checkbox"/> Yes <input type="checkbox"/> No Has the use of acute migraine medications (e.g., NSAIDs, triptans) decreased since the start of CGRP inhibitor therapy? <input type="checkbox"/> Yes <input type="checkbox"/> No Will the patient be initiating botulinum toxin headache prophylaxis after starting the requested agent? <input type="checkbox"/> Yes <input type="checkbox"/> No Does the patient continue to be monitored for medication overuse headache (MOH)? <input type="checkbox"/> Yes <input type="checkbox"/> No					



Please note: All information below is required to process this request.
Mon-Fri: 6am to 6pm Eastern / Sat: 6am to 6pm Eastern



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Quantity limit requests:

What is the quantity requested per MONTH? _____

Previous therapies failed and/or therapies currently used in combination with the requested medication (*List ALL medications tried or authorization process will be delayed*): _____

Is the prescribed dose higher than the maximum dose recommendation in FDA-approved labeling (i.e., the package insert)? Yes No

If **Yes**, please provide documentation to support the safety and efficacy of the higher dose (such as evidence from practice guidelines or clinical trials from peer-reviewed medical literature).

*** Please note: Chart documentation of the above is required to be submitted along with this fax*

****Use of samples to initiate therapy does not meet step edit/ preauthorization criteria.****

Previous therapies will be verified through pharmacy paid claims or submitted chart notes.

Prescriber Signature: _____ **Date:** _____

Are there any other comments, diagnoses, symptoms, medications tried or failed, and/or any other information the physician feels is important to this review?

Please note: This request may be denied unless all required information is received.
For urgent or expedited requests please call 1-800-711-4555.
This form may be used for non-urgent requests and faxed to 1-800-527-0531.